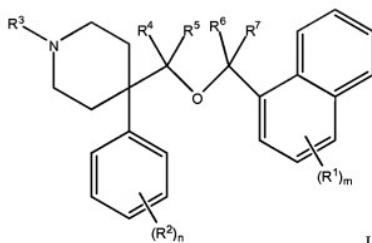


In the Claims:

This listing of claims will replace all prior versions, and listings of the claims in the application.

Please amend claims 1-5 and 8, cancel claims 7 and 9-13 without prejudice to their presentation in another application, and add new claims 14-29 as follows.

1. (currently amended) A compound in accord with formula I:



I

wherein:

R¹ at each occurrence is a moiety independently selected from CN, CF₃, OCF₃, OCHF₂, halogen, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, R^a, R^b, SR^a, NR^aR^f, CH₂NR^aR^f, OR^c, and CH₂OR^c, where m is selected from 0, 1, 2 or 3; wherein R^a, R^b, and R^c are independently at each occurrence selected from hydrogen, C₁₋₆alkyl, C(O)R^d, C(O)NHR^d and CO₂R^d, or R^a and R^b may together be (CH₂)_jG(CH₂)_k or G(CH₂)_j where G is oxygen, j is 1, 2, 3 or 4, k is 0, 1 or 2; where R^d at each occurrence is independently selected from C₁₋₆alkyl, and R^e and R^f are independently at each occurrence selected from hydrogen, C₁₋₆alkyl, C(O)R^d, C(O)NHR^d, and CO₂R^d;

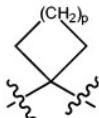
R² at each occurrence is a moiety independently selected from CN, CF₃, OCF₃, OCHF₂, halogen, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, R^a, R^b, SR^a, NR^aR^f, CH₂NR^aR^f, OR^c, and CH₂OR^c, where m is selected from 0, 1, 2 or 3; wherein R^a, R^b, and R^c are independently at each occurrence selected from hydrogen, C₁₋₆alkyl, C(O)R^d, C(O)NHR^d and CO₂R^d, or R^a and R^b may together be (CH₂)_jG(CH₂)_k or G(CH₂)_j where G is oxygen, j is 1, 2, 3 or 4, k is 0, 1 or

2; where R^d at each occurrence is independently selected from C₁₋₆alkyl, and R^e and R^f are independently at each occurrence selected from hydrogen, C₁₋₆alkyl, C(O)R^d, C(O)NHR^d, and CO₂R^d;

R³ is selected from hydrogen, C₁₋₆alkyl, C(O)-(CH₂)_q-NR⁸R⁹, (CH₂)_r-NR⁸R⁹, (CH₂)_q-O-D, (CH₂)_q-D and (CH₂)_q-CH=CH-D, wherein R⁸ and R⁹ are independently selected from hydrogen and C₁₋₆alkyl, q is selected from 1, 2 or 3, r is selected from 1, 2, 3 or 4 and D is selected from phenyl or indolyl which phenyl or indolyl may bear one or more substituents selected from halogen, C₁₋₆alkyl, C₁₋₆alkoxy and -O-(CH₂)_q-O-;

R⁴, R⁵, R⁶ and R⁷ at each occurrence are independently selected from hydrogen or C₁₋₆alkyl[[.]] ; or

independently, R⁴ and R⁵ together with the carbon to which they are attached and R⁶ and R⁷ together with the carbon to which they are attached form a moiety in accord with formula II,



wherein p is selected from 0, 1, 2, 3 or 4; or

in vivo hydrolysable precursors thereof, and or a pharmaceutically-acceptable salts salt thereof.

2. (currently amended) A compound according to Claim 1, wherein:

R¹ at each occurrence is independently selected from fluoro, cyano, C₁₋₆alkyl and C₁₋₆alkoxy and m is 1, 2 or 3;

R² at each occurrence is independently selected from halogen where n is 1 or 2, and

R³ is selected from hydrogen and C₁₋₆alkyl;

in vivo hydrolysable precursors thereof, and or a pharmaceutically-acceptable salts salt thereof.

3. (currently amended) A compound according to Claim 1, wherein:

R¹ at each occurrence is independently selected from fluoro, cyano, ethyl and methoxy

and m is 1, 2 or 3;

R² at each occurrence is independently selected from halogen where n is 1 or 2, and

R³ is selected from hydrogen and methyl;

in vivo-hydrolysable precursors thereof, and or a pharmaceutically-acceptable salts salt thereof.

4. (currently amended) A compound according to Claim 1, wherein R⁴, R⁵ and R⁶ are each hydrogen and R⁷ is methyl; *in vivo*-hydrolysable precursors thereof, and or a pharmaceutically-acceptable salts salt thereof.

5. (currently amended) A compound according to Claim 1, wherein:

R¹ at each occurrence is independently selected from fluoro, cyano, C₁₋₆alkyl and C₁₋₆alkoxy and m is 1, 2 or 3;

R² at each occurrence is independently selected from halogen where n is 1 or 2, and

R³ is selected from hydrogen, C₁₋₆alkyl, C(O)-(CH₂)_q-NR⁸R⁹, (CH₂)_r-NR⁸R⁹, (CH₂)_q-O-D, wherein R⁸ and R⁹ are independently selected from hydrogen, C₁₋₆alkyl and C₁₋₆alkoxy, q is 1, 2 or 3, r is 1, 2, 3 or 4 and D is selected from phenyl, indol-3-yl, indol-4-yl which phenyl may bear one or more substituents selected from fluoro, methyl, ethyl, methoxy, ethoxy or -O-(CH₂)₂-O- and which indolyl may bear one or more substituents selected from fluoro, methyl, ethyl, methoxy or ethoxy and ethoxy;

in vivo-hydrolysable precursors thereof, and or a pharmaceutically-acceptable salts salt thereof.

6. (original) A pharmaceutical composition comprising a compound according to Claim 1 together with at least one pharmaceutically-acceptable excipient or diluent.

7. (canceled).

8. (currently amended) A method of treating a disease condition wherein antagonism of NK₁ receptors in combination with SRI activity is beneficial which method comprises

comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 1 or ~~an in vivo hydrolysable precursor or~~ a pharmaceutically-acceptable salt thereof.

9-13. (canceled).

14. (new) A method of treating depression in cancer patients, depression in Parkinson's patients, postmyocardial infarction depression, subsyndromal symptomatic depression, depression in infertile women, pediatric depression, major depression, single episode depression, recurrent depression, child abuse induced depression, or post partum depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 1 or a pharmaceutically-acceptable salt thereof.

15. (new) A method of treating depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 1 or a pharmaceutically-acceptable salt thereof.

16. (new) A method of treating generalized anxiety disorder comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 1 or a pharmaceutically-acceptable salt thereof.

17. (new) A method of treating depression in cancer patients, depression in Parkinson's patients, postmyocardial infarction depression, subsyndromal symptomatic depression, depression in infertile women, pediatric depression, major depression, single episode depression, recurrent depression, child abuse induced depression, or post partum depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 2 or a pharmaceutically-acceptable salt thereof.

18. (new) A method of treating depression comprising administering to a warm-blooded

animal an effective amount of a compound according to Claim 2 or a pharmaceutically-acceptable salt thereof.

19. (new) A method of treating generalized anxiety disorder comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 2 or a pharmaceutically-acceptable salt thereof.

20. (new) A method of treating depression in cancer patients, depression in Parkinson's patients, postmyocardial infarction depression, subsyndromal symptomatic depression, depression in infertile women, pediatric depression, major depression, single episode depression, recurrent depression, child abuse induced depression, or post partum depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 3 or a pharmaceutically-acceptable salt thereof.

21. (new) A method of treating depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 3 or a pharmaceutically-acceptable salt thereof.

22. (new) A method of treating generalized anxiety disorder comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 3 or a pharmaceutically-acceptable salt thereof.

23. (new) A method of treating depression in cancer patients, depression in Parkinson's patients, postmyocardial infarction depression, subsyndromal symptomatic depression, depression in infertile women, pediatric depression, major depression, single episode depression, recurrent depression, child abuse induced depression, or post partum depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 4 or a pharmaceutically-acceptable salt thereof.

24. (new) A method of treating depression comprising administering to a warm-blooded

animal an effective amount of a compound according to Claim 4 or a pharmaceutically-acceptable salt thereof.

25. (new) A method of treating generalized anxiety disorder comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 4 or a pharmaceutically-acceptable salt thereof.

26. (new) A method of treating depression in cancer patients, depression in Parkinson's patients, postmyocardial infarction depression, subsyndromal symptomatic depression, depression in infertile women, pediatric depression, major depression, single episode depression, recurrent depression, child abuse induced depression, or post partum depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 5 or a pharmaceutically-acceptable salt thereof.

27. (new) A method of treating depression comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 5 or a pharmaceutically-acceptable salt thereof.

28. (new) A method of treating generalized anxiety disorder comprising administering to a warm-blooded animal an effective amount of a compound according to Claim 5 or a pharmaceutically-acceptable salt thereof.

29. (new) A method of treating depression in cancer patients, depression in Parkinson's patients, postmyocardial infarction depression, subsyndromal symptomatic depression, depression in infertile women, pediatric depression, major depression, single episode depression, recurrent depression, child abuse induced depression, or post partum depression comprising administering to a warm-blooded animal an effective amount of a compound selected from:

1-N-methyl-4-(3,4-dichlorophenyl)-4-((3-cyano-2-methoxynaphth-1-yl)methoxymethyl)piperidine;

4-(4-fluorophenyl)-4-[(3-cyano-2,4-dimethoxynaphth-1-yl)methoxymethyl]piperidine;
4-(4-fluorophenyl)-4-[(3-cyano-2-methoxynaphth-1-yl)methoxymethyl]piperidine;
{2-[4-(3-cyano-2-methoxy-naphthalen-1-yl-methoxymethyl)-4-(4-fluorophenyl)-
piperidin-1-yl]-2-oxo-ethyl}-methyl-carbamic acid tert-butyl ester;
4-[4-(4-fluorophenyl)-1-(2-methylamino-acetyl)-piperidin-4-ylmethoxymethyl]-3-
methoxy-naphthalene-2-carbonitrile;
4-{4-(4-fluorophenyl)-1-[2-(1H-indol-3-yl)-ethyl]-piperidin-4-ylmethoxymethyl}-3-
methoxy-naphthalene-2-carbonitrile;
1-N-methyl-4-(3,4-dichlorophenyl)-4-((3-cyanonaphth-1-yl)methoxymethyl)piperidine;
1-N-methyl-4-(3,4-dichlorophenyl)-4-((3-cyano-2,4-dimethoxynaphth-1-
yl)methoxymethyl)piperidine;
1-N-methyl-4-(4-fluorophenyl)-4-((3-cyanonaphth-1-yl)methoxymethyl)piperidine;
4-{{[4-fluoro-1-naphthyl)methoxy]methyl}-4-(4-fluorophenyl)-1-methylpiperidine;
4-(4-chlorophenyl)-4-[(3-cyano-2-methoxynaphth-1-yl)methoxymethyl]piperidine;
1-N-methyl-4-phenyl-4-((3-cyanonaphth-1-yl)methoxymethyl)piperidine;
1-N-methyl-4-(4-fluorophenyl)-4-((naphtha-1-yl)methoxymethyl)piperidine;
1-methyl-4-(1-naphthalen-1-yl-ethoxymethyl)-4-phenylpiperidine;
1-N-methyl-4-(4-fluorophenyl)-4-(3-cyano-1-naphthalen-1-yl-ethoxymethyl)piperidine;
and
1-N-methyl-4-(4-fluorophenyl)-4-(3-cyano-2,4-dimethoxy-1-naphthalen-1-yl-
ethoxymethyl)piperidine; or
a pharmaceutically-acceptable salt thereof.